In the claims:

1. (Original) An intraorally rapidly disintegrating tablet, characterized by being produced by tabletting cores coated with a pharmaceutical disintegrating agent, wherein the core is a granule containing a water-soluble medicament or containing a medicament and a sugar.

2. (Original) The intraorally rapidly disintegrating tablet according to claim 1, wherein the pharmaceutical disintegrating agent is crystalline cellulose, low-substituted hydroxypropyl cellulose, carboxymethyl cellulose, calcium carboxymethyl cellulose, crospovidone; or starch represented by potato starch, wheat starch, corn starch, rice starch, hydroxypropyl starch, sodium carboxymethyl starch, and partial-pregelatinized starch.

3. (Currently Amended) The intraorally rapidly disintegrating tablet according to claim 1 ~~or 2~~, wherein the sugar is selected from the group consisting of sugar alcohol represented by mannitol, xylitol, sorbitol, erythritol, maltitol and maltose; lactose, sucrose, glucose, and oligosaccharide.

4. (Currently Amended) The intraorally rapidly disintegrating tablet according to ~~any one of claims 1 to 3~~claim 1, wherein the average particle diameter of the coated granules is in the range of 20 to 1000 μm.

5. (Currently Amended) The intraorally rapidly disintegrating tablet according to ~~any one of claims 1 to 4~~claim 1, wherein the thickness of the tablet is in the range of 1 to 10 mm.